

Table 11.2 Principal characteristics of carcinomas and sarcomas

Feature	Carcinoma	Sarcoma
Origin	Epithelium	Connective tissues
Behaviour	Metastasis	Metastasis
Frequency	Common	Relatively rare
Preferred route of metastasis	Lymph	Blood
In situ phase	Yes	No
Age group	Usually over 50 years	Usually before 50 years

Although some general differences exist between the main groups of malignant tumours (Table 11.2), individual lesions have to be categorised more precisely both in clinical practice and for epidemiological purposes. It is inadequate to label the patient's tumour as merely having an epithelial or connective tissue origin; efforts must be made to determine the precise cell type. The classification of individual tumours is vitally important. Thorough histological examination of the tumour, sometimes using special techniques like genetic analysis and immunocytochemistry, detects subtle features that betray its provenance.

Histological grade (degree of differentiation)

The extent to which the tumour resembles histologically its cell or tissue of origin determines the tumour grade (Fig. 11.11) or degree of differentiation. Benign tumours are not usually further classified in this way because they nearly always closely resemble their parent tissue and grading the degree of differentiation offers no further clinical benefit in terms of choosing the most appropriate treatment. However, the degree of differentiation of malignant tumours is clinically useful both because it correlates strongly with patient survival (prognosis), and because it often indicates the most appropriate treatment. Thus, malignant tumours are usually graded either as well, moderately or poorly differentiated, or numerically, often by strict criteria, as grade 1, grade 2 or grade 3.

A well-differentiated tumour more closely resembles the parent tissue than does a poorly differentiated tumour, while moderately differentiated tumours are intermediate between these two extremes. Poorly differentiated tumours are more aggressive than well-differentiated tumours.

A few tumours are so poorly differentiated that they lack easily recognisable histogenetic features. There may even be great difficulty in deciding whether they are carcinomas or lymphomas, for example, although immunocytochemistry and genetic analysis often enable a distinction to be made. Tumours defying precise histogenetic classification are often referred to as 'anaplastic', or by some purely descriptive term such as 'spindle cell' or 'small round cell' tumour. Fortunately, advances



Benign and malignant tumours growing on skin, lower wall, showing the principal differences.

able prognosis and mortality associated with most may be due to

and destruction of adjacent tissue (secondary tumours (metastases) are observed in

of these (e.g. malignant tumour of the colon

of a tumour (e.g. ACTH and ADH from

metastatic effects causing weight loss and

skin

Classification

The cell of origin determines histological resemblance to parent tissue allows a graded mode correlates with clinical behaviour

A specific cell of origin of an individual tumour by histopathological examination and name type. This is then incorporated in the name (e.g. squamous cell carcinoma). Classification includes numerous subdivisions, origins of origin are:

of cells
of tissues
of and haemopoietic organs.

General And Systemic Pathology 5th Edition

Asma Khalil,Anthony Griffiths



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Table of Contents General And Systemic Pathology 5th Edition

1. Understanding the eBook General And Systemic Pathology 5th Edition
 - The Rise of Digital Reading General And Systemic Pathology 5th Edition
 - Advantages of eBooks Over Traditional Books
2. Identifying General And Systemic Pathology 5th Edition
 - Exploring Different Genres
 - Considering Fiction vs. Non-Fiction
 - Determining Your Reading Goals
3. Choosing the Right eBook Platform
 - Popular eBook Platforms
 - Features to Look for in an General And Systemic Pathology 5th Edition
 - User-Friendly Interface
4. Exploring eBook Recommendations from General And Systemic Pathology 5th Edition

- Personalized Recommendations
- General And Systemic Pathology 5th Edition User Reviews and Ratings
- General And Systemic Pathology 5th Edition and Bestseller Lists
- 5. Accessing General And Systemic Pathology 5th Edition Free and Paid eBooks
 - General And Systemic Pathology 5th Edition Public Domain eBooks
 - General And Systemic Pathology 5th Edition eBook Subscription Services
 - General And Systemic Pathology 5th Edition Budget-Friendly Options
- 6. Navigating General And Systemic Pathology 5th Edition eBook Formats
 - ePub, PDF, MOBI, and More
 - General And Systemic Pathology 5th Edition Compatibility with Devices
 - General And Systemic Pathology 5th Edition Enhanced eBook Features
- 7. Enhancing Your Reading Experience
 - Adjustable Fonts and Text Sizes of General And Systemic Pathology 5th Edition
 - Highlighting and Note-Taking General And Systemic Pathology 5th Edition
 - Interactive Elements General And Systemic Pathology 5th Edition
- 8. Staying Engaged with General And Systemic Pathology 5th Edition
 - Joining Online Reading Communities
 - Participating in Virtual Book Clubs
 - Following Authors and Publishers General And Systemic Pathology 5th Edition
- 9. Balancing eBooks and Physical Books General And Systemic Pathology 5th Edition
 - Benefits of a Digital Library
 - Creating a Diverse Reading Collection General And Systemic Pathology 5th Edition
- 10. Overcoming Reading Challenges
 - Dealing with Digital Eye Strain
 - Minimizing Distractions
 - Managing Screen Time
- 11. Cultivating a Reading Routine General And Systemic Pathology 5th Edition
 - Setting Reading Goals General And Systemic Pathology 5th Edition
 - Carving Out Dedicated Reading Time
- 12. Sourcing Reliable Information of General And Systemic Pathology 5th Edition

- Fact-Checking eBook Content of General And Systemic Pathology 5th Edition
- Distinguishing Credible Sources

13. Promoting Lifelong Learning

- Utilizing eBooks for Skill Development
- Exploring Educational eBooks

14. Embracing eBook Trends

- Integration of Multimedia Elements
- Interactive and Gamified eBooks

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